

FORM PTO-1390,

U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE

**TRANSMITTAL LETTER TO THE UNITED STATES
DESIGNATED/ELECTED OFFICE (DO/EO/US)
CONCERNING A FILING UNDER 35 U.S.C. 371**

ATTORNEY DOCKET NUMBER IN-5542	U.S. APPLICATION NO. IF KNOWN (SEE 37 CFR 1.5) 10/009161	
INTERNATIONAL APPLICATION NO. PCT/EP00/06035	INTERNATIONAL FILING DATE 29. June 2000 (29.06.2000)	PRIORITY DATE CLAIMED 30. June 1999 (30.06.99)

TITLE OF INVENTION: **ELECTRODEPOSITION BATH WITH WATER-SOLUBLE POLYVINYL ALCOHOL (CO)POLYMERS**

APPLICATION(S) FOR DO/EO/US: **Hardy REUTER, Dagmar SCHEMSCHAT, Karl-Heinz GROSSE-BRINKHAUS, Ulrich HEIMANN, Walter JOUCK**

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

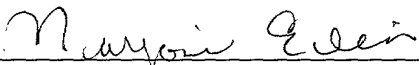
1. ☒ This is a FIRST submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a SECOND or SUBSEQUENT submission of items concerning a filing under 35 U.S.C. 371.
3. ☐ This express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371 (b) and PCT Articles 22 and 39(1).
4. ☐ A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date
5. ☒ A copy of the International Application as filed (35 U.S.C. 371(c)(2))
 - a. ☐ are transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☒ have been transmitted by the International Bureau.
 - c. ☐ is not required, as the application was filed in the United States Receiving Office (RO/US)
6. ☒ A translation of the International Application into English (35 U.S.C. 371(C)(2)).
7. ☒ Amendment to the claims of the International Application under PCT Article 19 (35 U.S.C.371(c)(3))
 - a. ☐ are transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☐ have been transmitted by the International Bureau.
 - c. ☐ have not been made; however, the time limit for making such amendments has NOT expired.
 - d. ☒ have not been made and will not be made
8. ☐ A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
9. ☒ An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).
10. ☐ A translation of the annex to the International Preliminary Examination Report under PCT Article 36


Items 11. to 16. below concern other document(s) or information included:

11. ☒ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
12. ☒ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included
13. ☒ A FIRST preliminary amendment.
☐ A SECOND or SUBSEQUENT preliminary amendment.
14. ☐ A substitute specification.
15. ☐ A Change of power of attorney and/or address letter.
16. ☒ Other items or information:

A copy of the cover sheet from the PCT Published Application
A copy of the cover sheet from the Priority Document

I hereby certify that the attached correspondence is being deposited with the United States Postal Service in an envelope as "Express Mail Post Office to Addressee" Mailing Label No. **ET332278465US** addressed to the Commissioner for Patents, Washington, 2900 Crystal Drive, Arlington 22202-3513 on December 5, 2001.


Marjorie Ellis

U.S. APPLICATION NO. 10/009161		INTERNATIONAL APPLICATION NO. PCT/EP00/06035		ATTORNEY'S DOCKET NUMBER IN-5542	
17. <input checked="" type="checkbox"/> The following fees are submitted				CALCULATIONS PTO USE ONLY	
Basic National Fee (37 CFR 1.492(a)(1)-(5)): Neither international preliminary examination fee (37 CFR 1.482) Nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO and International Search Report not prepared by the EPO or JPO.....				\$970.00	
International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International Search Report prepared by the EPO or JPO.....				\$890.00	
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ENTER APPROPRIATE BASIC FEE AMOUNT =				\$890.00	
Surcharge of \$130.00 for furnishing the oath or declaration later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(e)).				\$	
Claims	Number Filed	Number Extra	Rate		
Total Claims	16 - 20 =	0	X \$18.00	\$	
Independent claims	02 - 03 =	0	X \$80.00	\$	
Multiple dependent claims(s) (if applicable)			+ \$270.00	\$	
TOTAL OF ABOVE CALCULATION =				\$890.00	
Reduction by 1/2 for filing by small entity, if applicable. Verified Small Entity statement must also be filed. (Note 37 CFR 1.9, 1.27, 1.28).				\$	
SUBTOTAL =				\$890.00	
Processing fee of \$130.00 for furnishing the English translation later the <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(f)).				\$	
TOTAL NATIONAL FEE =				\$890.00	
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property +				\$	
TOTAL FEES ENCLOSED =				\$ 40.00	
				Amount to be:	\$
				refunded	
				Charged	\$930.00
a. <input type="checkbox"/> A check in the amount of \$_____ to cover the above fees is enclosed. b. <input checked="" type="checkbox"/> Please charge my Deposit Account No. <u>23-3425</u> in the amount of <u>\$930.00</u> to cover the above fees A triplicate copy of this sheet is enclosed. c. <input checked="" type="checkbox"/> The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. 23-3425. A triplicate copy of this sheet is enclosed.					
NOTE: Where an appropriate time limit under 37 CFR 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.					
SEND ALL CORRESPONDENCE TO: BASF CORPORATION Patent Department 26701 Telegraph Road Southfield, Michigan 48034-2442 (248) 948-2355 Customer No. 26922			 _____ SIGNATURE Michael F. Morgan _____ Name 42,906 _____ REGISTRATION NUMBER		

10/009161

JC10 Rec'd PCT/PTO 0 5 DEC 2001

PATENT

(Practitioner's Docket No. IN-5542)

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of

Hardy REUTER,
Dagmar SCHEMSCHAT,
Karl-Heinz GROSSE-BRINKHAUS,
Ulrich HEIMANN,
Walter JOUCK

Serial No.: This application is a National
Phase of Patent Application PCT/EP00/06035
filed on 29 June 2000.


Filed: December 5, 2001

For: ELECTRODEPOSITION BATH WITH
WATER-SOLUBLE POLYVINYL
ALCOHOL (CO)POLYMERS

Group Art Unit: Not Assigned

Examiner: Not Assigned

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Label No. **ET332278465US** addressed to the Box Patent
Application, Commissioner for Patents, 2900 Crystal Drive,
Arlington, VA 22202-3513 on December 5, 2001.


Marjorie Ellis

PRELIMINARY AMENDMENT UNDER 37 CFR § 1.115

BOX PATENT APPLICATION
Commissioner for Patents
Washington, D.C. 20231

Dear Sir:

This preliminary amendment is submitted with the application for entry into the U.S.
National Phase under Chapter II. This application is based on **PCT/EP00/06035** filed on 29
June 2000.

In connection with the filing of this National Phase application, please make the
following preliminary amendments.

IN THE SPECIFICATION

After the title, please insert --This application is a National Phase Application of
Patent Application PCT/EP00/06035 filed on 29 June 2000--

IN THE CLAIMS:

Please cancel claims 1-7 without prejudice or disclaimer.

Please add new claims 8-23.

8. (New) An aqueous electrodeposition bath comprising
- (A) a binder, wherein the binder is depositable cathodically or anodically,
 - (B) a dissolved polyvinyl alcohol (co)polymer comprising units of the following structure (I) $[-C(R^1)_2-C(R^1)(OH)-]$, wherein each R^1 in the structure is independently at least one of hydrogen, an alkyl, a substituted alkyl, a cycloalkyl, a substituted cycloalkyl, alkylcycloalkyl, substituted alkylcycloalkyl, cycloalkylalkyl, substituted cycloalkylalkyl, aryl, substituted aryl, alkylaryl, substituted alkylaryl, cycloalkylaryl, substituted cycloalkylaryl, arylalkyl, substituted arylalkyl, arylcycloalkyl, and substituted arylcycloalkyl, and
 - (C) optionally at least one of a crosslinking agent and a coatings additive.
9. (New) The aqueous electrodeposition bath of claim 8, wherein at least one R^1 in the structure is not hydrogen.
10. (New) The aqueous electrodeposition bath of claim 8, wherein the polyvinyl alcohol (co)polymer comprises a reaction product of structure (I) with at least one of a structure (II) $[-C(R^1)_2-C(R^1)(OC(O)R^2)-]$, wherein R^1 is as previously defined, and R^2 is a C_1 - C_{10} alkyl; a (meth)acrylic acid ester substantially free from acid groups; a monomer that carry at least one hydroxyl group per molecule and that are substantially free from acid groups; a monomer that carry per molecule at least one acid group that can be converted into a corresponding acid anion group; a vinyl ester of a C_5 - C_{18} alpha-branched monocarboxylic acid; a cyclic olefin, an acyclic olefin; (meth)acrylamide; a monomer containing an epoxide group; a vinylaromatic hydrocarbon; a nitrile; a vinyl monomer; and an allyl monomer.
11. (New) The aqueous electrodeposition bath of claim 8, wherein the polyvinyl alcohol (co)polymer is a copolymer of vinyl alcohol and at least one ethylenically unsaturated monomer.

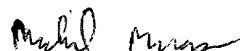
12. (New) The aqueous electrodeposition bath of claim 8, wherein the polyvinyl alcohol (co)polymer has a vinyl alcohol fraction of from 50 to 99.9 mol%.
13. (New) The aqueous electrodeposition bath of claim 8, wherein the weight average molecular mass of the polyvinyl alcohol (co)polymer is from 10,000 to 500,000 daltons.
14. (New) The aqueous electrodeposition bath of claim 8, wherein the polyvinyl alcohol (co)polymer is present in the electrodeposition bath in an amount from 2 to 10,000 ppm based on total weight of the electrodeposition bath.
15. (New) The aqueous electrodeposition bath of claim 8, wherein the coatings additive is at least one of an organic pigment, an inorganic pigment, an anticorrosion pigment, a filler, a free-radical scavenger, an organic corrosion inhibitor, a crosslinking catalyst, a slip additive, a polymerization inhibitor, a defoamer, an emulsifier, a wetting agent, an adhesion promoter, a leveling agent, a film-formation auxiliary, a flame retardant, an organic solvent, a reactive diluent that can participate in thermal crosslinking, and an anticrater agent.
16. (New) A method for coating an electrically conductive substrate, comprising
- (1) dipping the electrically conductive substrate into an electrodeposition bath as claimed in claim 8,
 - (2) connecting the substrate as one of the cathode or anode,
 - (3) applying a current to the substrate to deposit a film on the substrate,
 - (4) removing the substrate with the deposited film from the electrodeposition bath,
 - (5) baking the deposited coating film, and,
 - (6) optionally, following step (5), one of:
 - i) applying and baking a primer-surfacer, a stonechip protectant material, and a solid-color topcoat material, and
 - ii) applying and baking a basecoat material and a clearcoat material.
17. (New) The method of claim 16, wherein at least one R^1 in the structure is not hydrogen.

18. (New) The method of claim 16, wherein the polyvinyl alcohol (co)polymer comprises a reaction product of structure (I) with at least one of a structure (II) $[-C(R^1)_2-C(R^1)(OC(O)R^2)-]$, wherein R^1 is as previously defined, and R^2 is a C_1 - C_{10} alkyl; a (meth)acrylic acid ester substantially free from acid groups; a monomer that carry at least one hydroxyl group per molecule and that are substantially free from acid groups; a monomer that carry per molecule at least one acid group that can be converted into a corresponding acid anion group; a vinyl ester of a C_5 - C_{18} alpha-branched monocarboxylic acid; a cyclic olefin, an acyclic olefin; (meth)acrylamide; a monomer containing an epoxide group; a vinylaromatic hydrocarbon; a nitrile; a vinyl monomer; and an allyl monomer.
19. (New) The method of claim 16, wherein the polyvinyl alcohol (co)polymer is a copolymer of vinyl alcohol and at least one ethylenically unsaturated monomer.
20. (New) The method of claim 16, wherein the polyvinyl alcohol (co)polymer has a vinyl alcohol fraction of from 50 to 99.9 mol%.
21. (New) The method of claim 16, wherein the weight average molecular mass of the polyvinyl alcohol (co)polymer is from 10,000 to 500,000 daltons.
22. (New) The method of claim 16, wherein the polyvinyl alcohol (co)polymer is present in the electrodeposition bath in an amount from 2 to 10,000 ppm based on total weight of the electrodeposition bath.
23. (New) The method of claim 16, wherein the coatings additive is at least one of an organic pigment, an inorganic pigment, an anticorrosion pigment, a filler, a free-radical scavenger, an organic corrosion inhibitor, a crosslinking catalyst, a slip additive, a polymerization inhibitor, a defoamer, an emulsifier, a wetting agent, an adhesion promoter, a leveling agent, a film-formation auxiliary, a flame retardant, an organic solvent, a reactive diluent that can participate in thermal crosslinking, and an anticrater agent.

REMARKS

Upon entry of the present amendment claims 8-23 are pending in the application. New claims 8-23 add no new matter, as these claims contain subject matter from canceled claims 1-7 and the specification. There are a total of 16 claims and 1 independent claim pending in the application. Applicants respectfully request entry of the preliminary amendment.

Respectfully Submitted,



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PAT 99184 PCT
BASF Coatings AG

06.21.2000

**Electrodeposition bath comprising water-soluble
polyvinyl alcohol (co)polymers**

Description

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The invention relates to a novel use of water-soluble polyvinyl alcohol (co)polymers, to an electrodeposition bath comprising polyvinyl alcohol (co)polymers, and to coated substrates produced using same.

10

Electrodeposition coating is a well-known method of coating the surface of electrically conducting articles (compare, for example, Glasurit Handbuch Lacke und Farben, Curt R. Vincentz Verlag, Hanover, 1984, pages 374 to 384 and pages 457 to 462, and also DE-A-35 18 732, DE-A-35 18 770, EP-A-0 040 090, EP-A-0 012 463, EP-A-0 259 181, EP-A-0 433 783 and EP-A-0 262 069). The method is used to coat objects made of metal, especially for the priming of automobile bodies, or else to coat conductive plastics.

20

The coating materials used in electrodeposition coating generally comprise amino or carboxyl-containing synthetic resin binders, with dispersibility in water being achieved by neutralization of the amino or carboxyl groups. The electrodeposition coating materials may further include special grinding resins and possibly further constituents not dispersible in

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water, such as polymers, plasticizers, pigments, fillers, additives, and auxiliaries. The crosslinking agents used in the electrodeposition coating materials either are not dispersible in water or may be water-
5 dispersible, with the electrodeposition coating materials being externally crosslinking or else self-crosslinking, or being curable with condensation.

Modification to the binders, selection of the
10 crosslinkers, and variation of the composition of the ingredients of the electrodeposition coating material influence the properties of the coating, such as corrosion protection, adhesion, and leveling, for example. For instance, there have been disclosures in
15 particular of electrodeposition coating materials where by adding polymer microparticles or suspended and/or dispersed polymer powders the intention is to exert a favorable influence on corrosion protection, especially at edges, [lacuna] on leveling.

20

For instance, EP-A-0 259 181 recommends remedying the increased susceptibility to corrosion which is observed at edges of the coated substrate and is caused by a paint film of insufficient thickness by adding polymer
25 microgels, possible ingredients of such microgels being, for example, poly(meth)acrylate copolymers in combination with ethylenically unsaturated vinyl compounds.

Microgel dispersions which are based on epoxy-amine adducts and can be added subsequently are notable for their high compatibility and efficacy as edge protection additives, as described in EP 0 626 000.

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DE-B-26 50 611, EP-A-0 052 831, DE-A-39 40 782,
EP-A-0 433 783, SU-A-436890, JP-A-53094346,
JP-A-79028410 and JP-A-0624820 describe

10 electrodeposition coating compositions with suspendable
or dispersible polymer powders which are predominantly
free from ionic groups, are able to melt on baking if
desired, and are uncrosslinked or crosslinked, said
coating compositions further comprising the water-
dispersible synthetic resins that are typical of
15 electrodeposition coatings. The particle sizes of such
polymer powders may considerably exceed the particle
sizes of the water-dispersible synthetic resins of
known electrodeposition coating materials: the average
particle diameter in JP-A-0624820 is from 1 to 50
20 micrometers and in DE-A-39 40 782 or EP-A-0 433 783 is
from 0.1 to 100 micrometers.

In many cases, the addition of the polymer particles
described in EP-A-0 259 181, DE-B-26 50 611,
25 EP-A-0 052 831, EP-A-0 433 783, SU-A-436890,
JP-A-53094346, JP-A-79028410 and JP-A-0624820 to
aqueous electrodeposition coating materials leads to an
improvement in edge coverage. On the other hand,
despite the improved edge coverage, the corrosion

protection afforded by the deposited electrodeposition coating films, especially at the edges, is inadequate.

Disadvantageous side effects of adding polymer powders include a deterioration in the throwing power of the electrodeposition coating materials and in adhesion to the substrate and/or to subsequent coatings, such as paint films applied subsequently or PVC underbody protection, impairment of the mechanical properties, such as flexibility, stretchability, fracture strength and impact strength, poorer flow properties, and a drastic deterioration in leveling.

A furthermore [sic] key disadvantage of the aqueous and nonaqueous formulations described in the patents EP-A-0 259 181, DE-B-26 50 611, EP-A-0 052 831, EP-A-0 433 783, SU-A-436890, JP-A-53094346, JP-A-79028410, JP-A-0624820, SU-A-661637, SU-A-998592 and SU-A-310952 is the inadequate stability of the coating materials, which tend toward sedimentation. In aqueous electrodeposition coating materials, this may result in massive coverage of the ultrafiltration membrane with coarse polymer particles.

The stability disadvantages of the coating materials are alleviated by incorporating copolymers having vinyl acetal, vinyl alcohol and ethylene units directly into the resins, and/or by grafting reaction, as described in DE 196 18 379.

In this case a fraction of more than 10% by weight of polymer resin is needed in order to achieve sufficient edge coverage.

The incorporation of polymer powder or microgels requires fractions in the percent range, with a deterioration - in some instances drastic - in leveling.

Significantly more effective, even at low concentrations such as 500 ppm, in the electrodeposition coating material are water-soluble cellulose ethers, such as hydroxyethylcellulose (EP 0 640 700). The activity does not last, since the polymer degrades.

Polyvinyl alcohols are used multifariously in coating materials, in particular as suspension stabilizers for the polymerization of vinyl monomers. Whereas the use of polyvinyl alcohols as complexing agents and suspension stabilizers in the pretreatment of iron, steel, zinc and aluminum sheets, in combination with chromates and/or fluorine compounds, is known (J 73008702, WO 9627034), especially the electrophoretic deposition of metal suspensions, such as aluminum (SU 738334, J-A-111201), metal oxide suspensions, such as of chromium, aluminum, titanium and zirconium oxides (J-A-111201, SU 493817), metal salt suspensions, such as of lead, zinc or copper salts (SU 436890, SU 511392, SU 054452, WO 9208168), and also

direct deposition of metals, such as lead (SU 321265),
the direct use in electrodeposition coating materials
is restricted to subsequent treatment of the deposited
film by contact with an aqueous polyvinyl alcohol
5 solution followed by baking. This subsequent treatment
achieves a flatting effect (JP 56044799) or reduces
surface defects, such as craters (DE 4303787).

Against this background, the technical problem on which
10 the invention is based is that of specifying an
electrodeposition coating bath that gives coatings
which meet all of the requirements in respect of edge
protection and contamination resistance, especially to
oils, said coatings simultaneously being uncomplicated
15 to prepare and having long-term stability.

To solve this technical problem, the invention teaches
the use of a water-soluble polyvinyl alcohol
(co)polymer or of a mixture of polyvinyl alcohol
20 (co)polymers as an additive in aqueous
electrodeposition coating baths.

Aqueous electrodeposition coating baths contain little
if any organic solvent.

25

The expression water-soluble means a true solution
process in water and not a dispersion of particulate
units at a supermolecular level. Preferably, the
polyvinyl alcohol (co)polymer is prepared as an

additive in aqueous solution, where appropriate with customary coatings additaments, and the aqueous solution is added to the electrodeposition bath. The expression "additive" defines the presence of the
5 polyvinyl alcohol (co)polymer as a molecularly independent unit in the electrodeposition bath and in particular not as a component incorporated reactively into a binder, resin or the like. This definition does not of course rule out in a deposited coating the
10 polyvinyl alcohol (co)polymer being incorporated reactively into the other ingredients of the deposited coating.

In the context of the present invention, the term
15 polyvinyl alcohol (co)polymer refers to a random copolymer or block copolymer comprising polymer building blocks of the general formula I, or a homopolymer consisting of polymer building blocks of the general formula I, the polyvinyl alcohol copolymers
20 being of advantage in accordance with the invention and therefore being employed with preference.



25 In the polyvinyl alcohol (co)polymers for use in accordance with the invention the polymer building blocks I may be linked head to head or head to tail. Advantageously, by far the predominant proportion of

the polymer building blocks I are linked head to tail.

In the general formula I, the variable R^1 stands for hydrogen atoms or for substituted or unsubstituted
5 alkyl, cycloalkyl, alkylcycloalkyl, cycloalkylalkyl, aryl, alkylaryl, cycloalkylaryl, arylalkyl or arylcycloalkyl radicals.

Examples of suitable alkyl radicals are methyl, ethyl,
10 propyl, isopropyl, n-butyl, isobutyl, tert-butyl, amyl, hexyl, and 2-ethylhexyl.

Examples of suitable cycloalkyl radicals are cyclobutyl, cyclopentyl, and cyclohexyl.

15 Examples of suitable alkylcycloalkyl radicals are methylenecyclohexane, ethylenecyclohexane, and propane-1,3-diylcyclohexane.

20 Examples of suitable cycloalkylalkyl radicals are 2-, 3- and 4-methyl-, -ethyl-, -propyl-, and -butylcyclohex-1-yl.

Examples of suitable aryl radicals are phenyl,
25 naphthyl, and biphenyl.

Examples of suitable alkylaryl radicals are benzyl-
[sic], ethylene- and propane-1,3-diyl-benzene.

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Examples of suitable cycloalkylaryl radicals are 2-, 3-, and 4-phenylcyclohex-1-yl.

Examples of suitable arylalkyl radicals are 2-, 3- and 4-methyl-, -ethyl-, -propyl-, and -butylphen-1-yl.

Examples of suitable arylcycloalkyl radicals are 2-, 3-, and 4-cyclohexylphen-1-yl.

10 The above-described radicals R^1 may be substituted. Electron-withdrawing or electron-donating atoms or organic radicals may be used for this purpose.

Examples of suitable substituents are halogen atoms, especially chlorine or fluorine, nitrile groups, nitro groups, partly or fully halogenated, especially chlorinated and/or fluorinated, alkyl, cycloalkyl, alkylcycloalkyl, cycloalkylalkyl, aryl, alkylaryl, cycloalkylaryl, arylalkyl and arylcycloalkyl radicals, including those exemplified above, especially tert-butyl; aryloxy, alkyloxy and cycloalkyloxy radicals, especially phenoxy, naphthoxy, methoxy, ethoxy, propoxy, butyloxy or cyclohexyloxy; arylthio, alkylthio and cycloalkylthio radicals, especially phenylthio, naphthylthio, methylthio, ethylthio, propylthio, butylthio or cyclohexylthio; hydroxyl groups; and/or primary, secondary and/or tertiary amino groups, especially amino, N-methylamino, N-ethylamino, N-propylamino, N-phenylamino, N-cyclohexylamino,

N,N-dimethylamino, N,N-diethylamino, N,N-dipropylamino, N,N-diphenylamino, N,N-dicyclohexylamino, N-cyclohexyl-N-methylamino or N-ethyl-N-methylamino.

- 5 It is of advantage in accordance with the invention if the radicals R^1 comprise predominantly hydrogen atoms, i.e., if the other radicals R^1 are present only to a minor extent. In the context of the present inventions [sic], the term "minor extent" designates an extent
- 10 which advantageously varies and does not impair or even completely alter the profile of performance properties of the polyvinyl alcohol (co)polymers, especially their solubility in water. Particular advantages result if the radicals R^1 comprise exclusively hydrogen atoms,
- 15 i.e., if the polymer building blocks I are derived from the hypothetical polyvinyl alcohol. Accordingly, polyvinyl alcohol (co)polymers containing these polymer building blocks I are used with particular preference.
- 20 Besides the polymer building blocks I, the polyvinyl alcohol copolymers for use in accordance with the invention further comprise, in particular, polymer building blocks of the general formula II.



25

In the general formula II, the radicals R^1 have the definition indicated above, hydrogen atoms again being

of particular advantage and therefore being employed with particular preference. The radicals R^2 stand for alkyl radicals having from one to ten carbon atoms, preferably methyl, ethyl, propyl, isopropyl, n-butyl, isobutyl, tert-butyl, amyl, hexyl, or 2-ethylhexyl, with particular preference methyl. Accordingly, the particularly preferred polymer building blocks II are derived from vinyl acetate. The polymer building blocks II may be linked head to head or head to tail. Advantageously, by far the predominant proportion of the polymer building blocks II are linked head to tail.

The polyvinyl alcohol copolymers may further comprise customary and known ethylenically unsaturated monomers such as

- (meth)acrylic esters substantially free from acid groups,
- monomers which carry at least one hydroxyl group per molecule and are substantially free from acid groups, such as hydroxyalkyl esters of acrylic acid, methacrylic acid or another alpha,beta-olefinically unsaturated carboxylic acid which are derived from an alkylene glycol which is esterified with the acid or are obtainable by reacting the alpha,beta-olefinically unsaturated carboxylic acid with an alkylene oxide,

- monomers which carry per molecule at least one acid group which can be converted into the corresponding acid anion group,
- 5 - vinyl esters of alpha-branched monocarboxylic acids having from 5 to 18 carbon atoms in the molecule,
- reaction products of acrylic acid and/or
10 methacrylic acid with the glycidyl ester of an alpha-branched monocarboxylic acid having from 5 to 18 carbon atoms per molecule,
- cyclic and/or acyclic olefins such as ethylene,
15 propylene, but-1-ene, pent-1-ene, hex-1-ene, cyclohexene, cyclopentene, norbornene, butadiene, isoprene, cyclopentadiene and/or dicyclopentadiene, especially ethylene,
- 20 - (meth)acrylamides,
- monomers containing epoxide groups, such as the glycidyl esters of ethylenically unsaturated carboxylic acids,
- 25 - vinylaromatic hydrocarbons,
- nitriles,

- vinyl compounds, especially vinyl halides and/or vinylidene dihalides, N-vinylpyrrolidone or vinyl ethers,
- 5 - allyl compounds, especially allyl ethers and allyl esters.

Where these monomers are used, they are present only in a minor extent in the polyvinyl alcohol copolymers for use in accordance with the invention, this term being employed here again in the sense explained above. Of these monomers, the acyclic olefins, especially ethylene and propylene, in particular ethylene, offer particular advantages and are therefore used with preference where needed.

Advantageously, the polyvinyl alcohol (co)polymers for use in accordance with the invention have a degree of polymerization of from 100 to 20 000, preferably from 200 to 15 000, with particular preference from 300 to 12 000, and in particular from 400 to 10 000.

The amount of polymer building blocks I in the polyvinyl alcohol copolymers is advantageously from 50 to 99.9, with preference from 60 to 99.9, with particular preference from 70 to 99 and in particular from 80 to 99 mol%.

In the context of the present invention, the polyvinyl

alcohol copolymers which comprise the particularly advantageous polymer building blocks I and II offer very particular advantages and are therefore used with very particular preference in accordance with the invention. These polyvinyl alcohol copolymers are also referred to for short by those in the art as polyvinyl alcohols.

As is known, the polyvinyl alcohols are not accessible through direct polymerization processes but instead are prepared by way of polymer-analogous reactions by hydrolysis of polyvinyl acetate. Particularly advantageous, commercially customary polyvinyl alcohols have molecular weights of from 10 000 to 500 000 daltons, preferably from 15 000 to 320 000 daltons, and in particular from 20 0000 [sic] to 300 000 daltons. Especially advantageous, commercially customary polyvinyl alcohols have a degree of hydrolysis of from 98 to 99 or from 87 to 89 mol%.

20

The vinyl alcohol fraction may be determined, for example, indirectly by way of the ester number in accordance with DIN 53401, specifically by determining the remaining fraction of vinyl acetate following hydrolysis by means of the ester number.

25

The solubility of these polyvinyl alcohols in water may be varied within a wide range by subsequent polymer-analogous modification with aldehydes. As is known,

this reaction leads to the formation of cyclic acetals. Examples of suitable acetalized polyvinyl alcohols are known from the patent DE-A-196 18 379.

5 Surprisingly, the addition of the polyvinyl alcohol (co)polymer, especially polyvinyl alcohol, which is easy to prepare and may be added simply as an additive directly to the electrodeposition bath, achieves edge protection to satisfy every requirement, and very good
10 contamination resistance, especially to oil. The leveling is likewise outstanding. It has been found, furthermore, that only very small amounts of polyvinyl alcohol (co)polymer need be added, leading to a considerable cost advantage over the prior art's added
15 edge protection improvers.

In the context of the invention it is advantageous if the fraction of polyvinyl alcohol (co)polymers, especially polyvinyl alcohols, in the electrodeposition
20 bath is from 2 to 10 000 ppm, preferably from 20 to 5000 ppm, based in each case on the total weight of the electrodeposition bath. If the electrodeposition bath includes pigments (inorganic) in a fraction of more than 10%, based on the solids of the electrodeposition
25 bath, then it is generally sufficient to add an amount of from 2 to 3000, in particular 300 to 1500, ppm.

The use in accordance with the invention is advantageous in the context of all customary anodic or

cathodic electrodeposition baths.

These electrodeposition baths are aqueous coating materials having a solids content of in particular from 5 . 5 to 30% by weight.

The solids of the bath of the invention comprises

10 (A) customary and known binders which carry ionic groups or functional groups which can be converted into ionic groups, (a1), and also functional groups (a2) capable of chemical crosslinking, and are externally crosslinking and/or self-crosslinking, but in particular externally
15 crosslinking;

(B) if desired, crosslinking agents which carry complementary functional groups (b1) which are able to undergo chemical crosslinking reactions
20 with the functional groups (a2) and are employed mandatorily when the binders (A) are externally crosslinking;

(C) customary and known coatings additives, and
25

(D) the polyvinyl alcohol (co)polymers for use in accordance with the invention and described in detail above, especially polyvinyl alcohols.

Where the crosslinking agents (B) and/or their functional groups (b1) have already been incorporated into the binders (A), the term self-crosslinking is used.

5

Suitable complementary functional groups (a2) of the binders (A) include, preferably, thio, amino, hydroxyl, carbamate, allophanate, carboxyl, and/or (meth)acrylate groups, but especially hydroxyl groups, and

10 complementary functional groups (b1) include preferably anhydride, carboxyl, epoxy, blocked isocyanate, urethane, methylol, methylol ether, siloxane, amino, hydroxyl and/or beta-hydroxyalkylamide groups, but especially blocked isocyanate groups.

15

Examples of suitable ionic functional groups, or functional groups which can be converted into ionic groups, (a1), of the binders (A) are

20 (a11) functional groups which can be converted into cations by neutralizing agents and/or quaternizing agents, and/or cationic groups

or

25

(a12) functional groups which can be converted into anions by neutralizing agents, and/or anionic groups.

The binders (A) having functional groups (a11) are used in cathodically depositable electrodeposition coating (cathodic electrocoat) materials, while the binders (A) having functional groups (a12) are employed in anodic electrocoat materials.

Examples of suitable functional groups (a11) for use in accordance with the invention that can be converted into cations by neutralizing agents and/or quaternizing agents are primary, secondary or tertiary amino groups, secondary sulfide groups or tertiary phosphine groups, especially tertiary amino groups or secondary sulfide groups.

Examples of suitable cationic groups (a11) for use in accordance with the invention are primary, secondary, tertiary or [sic] tertiary sulfonium groups or quaternary phosphonium groups, preferably quaternary ammonium groups or quaternary ammonium groups [sic], tertiary sulfonium groups, but especially quaternary ammonium groups.

Examples of suitable functional groups (a12) for use in accordance with the invention that can be converted into anions by neutralizing agents are carboxylic acid, sulfonic acid or phosphonic acid groups, especially carboxylic acid groups.

Examples of suitable anionic groups (a12) for use in

accordance with the invention are carboxylate, sulfonate or phosphonate groups, especially carboxylate groups.

- 5 The groups (a11) or (a12) should be selected so as to rule out the possibility of disruptive reactions with the functional groups (a2) that are able to react with the crosslinking agents (B). The skilled worker will therefore be able to make the selection in a simple
10 manner on the basis of his or her knowledge of the art.

Examples of suitable neutralizing agents for functional groups (a11) convertible into cations are inorganic and organic acids such as sulfuric acid, hydrochloric acid,
15 phosphoric acid, formic acid, acetic acid, lactic acid, dimethylolpropionic acid or citric acid, especially formic acid, acetic acid or lactic acid.

Examples of suitable neutralizing agents for functional
20 groups (a12) convertible into anions are ammonia, ammonium salts, such as ammonium carbonate or ammonium hydrogen carbonate, for example, and also amines, such as trimethylamine, triethylamine, tributylamine, dimethylaniline, diethylaniline, triphenylamine,
25 dimethylethanolamine, diethylethanolamine, methyldiethanolamine, triethanolamine and the like.

The amount of neutralizing agent is generally is [sic] chosen so that from 1 to 100 equivalents, preferably

from 50 to 90 equivalents, of the functional groups (a11) or (a12) of the binder (b1) are neutralized.

Examples of suitable binders (A) for anodic electrocoat materials are known from the patent DE-A-28 24 418. They comprise preferably polyesters, epoxy resin esters, poly(meth)acrylates, maleate oils or polybutadiene oils having a weight average molecular weight of from 300 to 10 000 daltons and an acid number of from 35 to 300 mg KOH/g.

Examples of suitable cathodic electrocoat materials are known from the patents EP-A-0 082 291, EP-A-0 234 395, EP-A-0 227 975, EP-A-0 178 531, EP-A-333 327, EP-A-0 310 971, EP-A-0 456 270, US-A-3,922,253, EP-A-0 261 385, EP-A-0 245 786, DE-A-33 24 211, EP-A-0 414 199, and EP-A-476 514. They comprise preferably resins (A) containing primary, secondary, tertiary or quaternary amino or ammonium groups and/or tertiary sulfonium groups and having amine numbers of preferably between 20 and 250 mg KOH/g and a weight average molecular weight of preferably from 300 to 10 000 daltons. In particular, amino (meth)acrylate resins, amino [sic] epoxy resins, amino epoxy resins having terminal double bonds, amino epoxy resins having primary and/or secondary hydroxyl groups, amino polyurethane resins, amino-containing polybutadiene resins, or modified epoxy resin/carbon dioxide/amine reaction products [lacuna].

In accordance with the invention, cathodic electrocoat materials and the corresponding electrodeposition baths are used with preference.

5

The electrodeposition baths preferably comprise crosslinking agents (B).

Examples of suitable crosslinking agents (B) are
10 blocked organic polyisocyanates, especially blocked polyisocyanates known as paint polyisocyanates, containing blocked isocyanate groups attached to aliphatic, cycloaliphatic, araliphatic and/or aromatic moieties.

15

They are preferably prepared using polyisocyanates having from 2 to 5 isocyanate groups per molecule and having viscosities of from 100 to 10 000, preferably from 100 to 5000, and in particular from 100 to
20 2000 mPas (at 23°C). Moreover, the polyisocyanates may have been subjected to conventional hydrophilic or hydrophobic modification.

Examples of suitable polyisocyanates are described, for
25 example, in "Methoden der organischen Chemie", Houben-Weyl, Volume 14/2, 4th edition, Georg Thieme Verlag, Stuttgart 1963, pages 61 to 70, and by W. Siefken, Liebigs Annalen der Chemie, Volume 562, pages 75 to 136. Examples of those suitable include the isocyanato-

containing polyurethane prepolymers which can be prepared by reacting polyols with an excess of polyisocyanates and which are preferably of low viscosity.

5

Further examples of suitable polyisocyanates are polyisocyanates containing isocyanurate, biuret, allophanate, iminooxadiazinedione, urethane, urea and/or uretdione groups. Polyisocyanates containing urethane groups, for example, are obtained by reacting some of the isocyanate groups with polyols, such as trimethylolpropane and glycerol, for example. It is preferred to use aliphatic or cycloaliphatic polyisocyanates, especially hexamethylene diisocyanate, dimerized and trimerized hexamethylene diisocyanate, isophorone diisocyanate, 2-isocyanatopropylcyclohexyl isocyanate, dicyclohexylmethane 2,4'-diisocyanate, dicyclohexylmethane 4,4'-diisocyanate or 1,3-bis(isocyanatomethyl)cyclohexane (BIC), diisocyanates derived from dimer fatty acids, as sold under the commercial designation DDI 1410 by Henkel, 1,8-diisocyanato-4-isocyanatomethyloctane, 1,7-diisocyanato-4-isocyanatomethylheptane or 1-isocyanato-2-(3-isocyanatopropyl)-cyclohexane, or mixtures of these polyisocyanates.

25

Examples of suitable blocking agents for preparing the blocked polyisocyanates (B) are the blocking agents known from the US patent US-A-4,444,954, such as

- i) phenols such as phenol, cresol, xylenol, nitrophenol, chlorophenol, ethylphenol, t-butylphenol, hydroxybenzoic acid, esters of this acid, or 2,5-di-tert-butyl-4-hydroxytoluene;
- 5
- ii) lactams, such as ϵ -caprolactam, δ -valerolactam, γ -butyrolactam or β -propiolactam;
- 10
- iii) active methylenic compounds, such as diethyl malonate, dimethyl malonate, ethyl or methyl acetoacetate, or acetylacetone;
- 15
- iv) alcohols such as methanol, ethanol, n-propanol, isopropanol, n-butanol, isobutanol, t-butanol, n-amyl alcohol, t-amyl alcohol, lauryl alcohol, ethylene glycol monomethyl ether, ethylene glycol monoethyl ether, ethylene glycol monobutyl ether, diethylene glycol monomethyl ether, diethylene glycol monoethyl ether, propylene glycol monomethyl ether, methoxymethanol, glycolic acid, glycolic esters, lactic acid, lactic esters, methylolurea, methylolmelamine, diacetone alcohol, ethylenechlorohydrin, ethylenebromohydrin, 1,3-dichloro-2-propanol, 1,4-cyclohexyldimethanol or
- 20
- 25
- acetocyanohydrin;
- v) mercaptans such as butyl mercaptan, hexyl mercaptan, t-butyl mercaptan, t-dodecyl mercaptan, 2-mercaptobenzothiazole, thiophenol, methylthio-

phenol or ethylthiophenol;

- vi) acid amides such as acetoanilide, acetoanisidin-
amide, acrylamide, methacrylamide, acetamide,
5 stearamide or benzamide;
- vii) imides such as succinimide, phthalimide or
maleimide;
- 10 viii) amines such as diphenylamine, phenyl-naphthylamine,
xylidine, N-phenylxylidine, carbazole, aniline,
naphthylamine, butylamine, dibutylamine or butyl-
phenylamine;
- 15 ix) imidazoles such as imidazole or 2-ethylimidazole;
- x) ureas such as urea, thiourea, ethyleneurea,
ethylenethiourea or 1,3-diphenylurea;
- 20 xi) carbamates such as phenyl N-phenylcarbamate or
2-oxazolidone;
- xii) imines such as ethyleneimine;
- 25 xiii) oximes such as acetone oxime, formaldoxime,
acetaldoxime, acetoxime, methyl ethyl ketoxime,
diisobutyl ketoxime, diacetyl monoxime, benzo-
phenone oxime or chlorohexanone oximes;

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xiv) salts of sulfurous acid such as sodium bisulfite or potassium bisulfite;

xv) hydroxamic esters such as benzyl methacrylo-
5 hydroxamate (BMH) or allyl methacrylohydroxamate;
or

xvi) substituted pyrazoles, ketoximes imidazoles or triazoles; and also

10

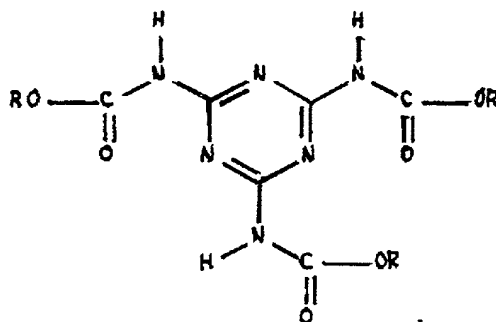
mixtures of these blocking agents, especially dimethylpyrazole and triazoles, malonic esters and acetoacetic esters or dimethylpyrazole and succinimide, or butyl diglycol and trimethylolpropane.

15

Further examples of suitable crosslinking agents (B) are all known aliphatic and/or cycloaliphatic and/or aromatic polyepoxides, based for example on bisphenol A or bisphenol F. Further suitable polyepoxides include,
20 for example, the polyepoxides obtainable commercially under the designations Epikote® from Shell, Denacol® from Nagase Chemicals Ltd., Japan, such as, for example Denacol EX-411 (pentaerythritol polyglycidyl ether), Denacol EX-321 (trimethylolpropane polyglycidyl ether),
25 Denacol EX-512 (polyglycerol polyglycidyl ether), and Denacol EX-521 (polyglycerol polyglycidyl ether).

As crosslinking agents (B) it is also possible to use tris(alkoxycarbonylamino)triazines (TACT) of the

general formula



5 Examples of suitable tris(alkoxycarbonylamino)triazines (B) are described in the patents US-A-4,939,213, US-A-5,084,541, and EP-A-0 624 577. Use is made in particular of the tris(methoxy-, tris(butoxy- and/or tris(2-ethylhexoxycarbonylamino)triazines.

10

The methyl butyl mixed esters, the butyl 2-ethylhexyl mixed esters, and the butyl esters are of advantage. They have the advantage over the straight methyl ester of better solubility in polymer melts, and also have
15 less of a tendency to crystallize out.

Further examples of suitable crosslinking agents (B) are amino resins, examples being melamine resins, guanamine resins, benzoguanamine resins or urea resins.

20 Also suitable are the customary and known amino resins some of whose methylol and/or methoxymethyl groups have been defunctionalized by means of carbamate or allophanate groups. Crosslinking agents of this kind are described in the patents US-A-4,710,542 and

EP-B-0 245 700 and also in the article by B. Singh and coworkers, "Carbamylmethyated Melamines, Novel Crosslinkers for the Coatings Industry" in Advanced Organic Coatings Science and Technology Series, 1991, 5 Volume 13, pages 193 to 207.

Further examples of suitable crosslinking agents (B) are beta-hydroxyalkylamides such as N,N,N',N'-tetrakis(2-hydroxyethyl)adipamide or N,N,N',N'-tetra- 10 kis(2-hydroxypropyl)adipamide.

Further examples of suitable crosslinking agents (B) are compounds containing on average at least two groups amenable to transesterification, examples being 15 reaction products of malonic diesters and polyisocyanates or of monoisocyanates with esters and partial esters of malonic acid with polyhydric alcohols, as described in the European patent EP-A-0 596 460.

20

The amount of the crosslinking agents (B) in the coating material or electrodeposition bath of the invention may vary widely and is guided in particular, firstly, by the functionality of the crosslinking 25 agents (B) and, secondly, by the number of crosslinking functional groups (a₂) which are present in the binder (A), and also by the target crosslinking density. The skilled worker is therefore able to determine the amount of the crosslinking agents (B) on the basis of

his or her general knowledge in the art, possibly with the aid of simple rangefinding experiments. Advantageously, the crosslinking agent (B) is present in the coating material of the invention in an amount
5 of from 5 to 60, with particular preference from 10 to 50, and in particular from 15 to 45% by weight, based in each case on the solids content of the coating material of the invention. It is further advisable here to choose the amounts of crosslinking agent (B) and
10 binder (A) such that in the coating material of the invention the ratio of functional groups (b1) in the crosslinking agent (B) to functional groups (a2) in the binder (A) is from 2:1 to 1:2, preferably from 1.5:1 to 1:1.5, with particular preference from 1.2:1 to 1:1.2,
15 and in particular from 1.1:1 to 1:1.1.

The coating material or electrodeposition bath of the invention may comprise customary coatings additives (C) in effective amounts. Examples of suitable additives
20 (C) are

- organic and/or inorganic pigments, anticorrosion pigments and/or fillers such as calcium sulfate, barium sulfate, silicates such as talc or kaolin,
25 silicas, oxides such as aluminum hydroxide or magnesium hydroxide, nanoparticles, organic fillers such as textile fibers, cellulose fibers, polyethylene fibers or woodflour, titanium dioxide, carbon black, iron oxide, zinc phosphate

or lead silicate; these additives may also be incorporated into the electrodeposition baths of the invention by way of pigment pastes, suitable grinding resins comprising the binders (A) described above;

- free-radical scavengers;

- organic corrosion inhibitors;

- crosslinking catalysts such as organic and inorganic salts and complexes of tin, lead, antimony, bismuth, iron or manganese, preferably organic salts and complexes of bismuth and of tin, especially bismuth lactate, ethylhexanoate or dimethylolpropionate and dibutyltin oxide or dibutyltin dilaurate;

- slip additives;

- polymerization inhibitors;

- defoamers;

- emulsifiers, especially nonionic emulsifiers such as alkoxylated alkanols and polyols, phenols and alkylphenols or anionic emulsifiers such as alkali metal salts or ammonium salts of alkanecarboxylic acids, alkanesulfonic acids, and sulfo acids of

alkoxylated alkanols and polyols, phenols and alkylphenols;

- 5 - wetting agents such as siloxanes, fluorine compounds, carboxylic monoesters, phosphoric esters, polyacrylic acids and their copolymers, or polyurethanes;
- 10 - adhesion promoters;
- leveling agents;
- film-formation auxiliaries such as cellulose derivatives;
- 15 - flame retardants;
- organic solvents;
- 20 - low molecular mass, oligomeric and high molecular mass reactive diluents which can participate in thermal crosslinkings, especially polyols such as tricyclodecanedimethanol, dendrimeric polyols, hyperbranched polyesters, polyols based on
- 25 metathesis oligomers or branched alkanes having more than eight carbon atoms in the molecule;
- anticrater agents;

Further examples of suitable coatings additives are described in the textbook "Lackadditive" [Additives for coatings] by Johan Bieleman, Wiley-VCH, Weinheim, New York, 1998.

5

Finally, the invention teaches a method of coating electrically conductive substrates, in which (1) the electrically conductive substrate is dipped into an electrodeposition bath in accordance with the information given above, (2) the substrate is connected as the cathode or anode, preferably as the cathode, (3) a film is deposited on the substrate by means of direct current, (4) the coated substrate is removed from the electrodeposition bath, (5) the deposited coating film is baked, and (6) optionally, following step (5), a primer-surfacer, a stonechip protectant material and a solid-color topcoat material, or alternatively a basecoat material and a clearcoat material, are applied and baked, the basecoat material and the clearcoat material preferably being applied and baked by the wet-on-wet technique.

Examples

1. Preparation of the crosslinking agents (B)

1.1 Preparation of the crosslinking agent (B1)

A reactor equipped with a stirrer, reflux condenser,

internal thermometer and inert gas inlet is charged under a nitrogen atmosphere with 10552 parts of isomers and more highly functional oligomers based on 4,4'-diphenylmethane diisocyanate having an NCO equivalent weight of 135 g/eq (Lupranat®, BASF/Germany; NCO functionality approximately 2.7; 2,2'- and 2,4'-diphenylmethane diisocyanate content less than 5%). 18 parts of dibutyltin dilaurate are added and 9498 parts of butyl diglycol are added dropwise at a rate such that the product temperature remains below 60°C. Cooling may be necessary. After the end of the addition, the temperature is held at 60°C for a further 60 minutes and an NCO equivalent weight of 1120 g/eq is found (based on solids fractions). Following dilution in 7768 parts of methyl isobutyl ketone, 933 parts of melted trimethylolpropane are added at a rate such that the product temperature does not exceed 100°C. After the end of the addition, reaction is continued for 60 minutes more. On subsequent checking, NCO groups are no longer detectable. Cooling to 65°C is accompanied by dilution with 965 parts of n-butanol and 267 parts of methyl isobutyl ketone.

The solids content is 70.1% (1 h at 130°C).

1.2 Preparation of the crosslinking agent (B2)

A reactor equipped with a stirrer, reflux condenser, internal thermometer and inert gas inlet is charged

under a nitrogen atmosphere with 12208 parts of isomers and more highly functional oligomers based on 4,4'-diphenylmethane diisocyanate having an NCO equivalent weight of 135 g/eq (Lupranat®, BASF/Germany; 5 NCO functionality approximately 2.7; 2,2'- and 2,4'-diphenylmethane diisocyanate content less than 5%). 8 parts of dibutyltin dilaurate are added and 10499 parts of butyl diglycol are added dropwise at a rate such that the product temperature remains below 10 60°C. Cooling may be necessary. After the end of the addition, the temperature is held at 60°C for a further 60 minutes and an NCO equivalent weight of 887 g/eq is found (based on solids fractions). Following dilution in 4500 parts of methyl isobutyl ketone, 1293 parts of 15 melted trimethylolpropane are added at a rate such that the product temperature does not exceed 100°C. After the end of the addition, reaction is continued for 60 minutes more. On subsequent checking, NCO groups are no longer detectable. Cooling to 65°C is accompanied by 20 dilution with 599 parts of n-butanol and 893 parts of methyl isobutyl ketone.

The solids content is 80.5% (1 h at 130°C).

25 **2. Preparation of the precursor (solution of diethylenetriamine diketimine in methyl isobutyl ketone)**

From a 70 percent strength by weight solution of

diethylenetriamine in methyl isobutyl ketone, the water of reaction is removed azeotropically at 110-140°C. This is followed by dilution with methyl isobutyl ketone until the solution has an amine equivalent weight of 127.

3. Preparation of aqueous dispersions containing cathodically depositable resins (A) and a crosslinking agent (B)

3.1 Preparation of the aqueous binder dispersion (A/B1)

In a reactor equipped with a stirrer, reflux condenser, internal thermometer and inert gas inlet, 6150 parts of epoxy resin based on bisphenol A having an epoxy equivalent weight (EEW) of 188 are heated to 125°C under a nitrogen atmosphere together with 1400 parts of bisphenol A, 335 parts of dodecylphenol, 470 parts of p-cresol and 441 parts of xylene and the mixture is stirred for 10 minutes. It is subsequently heated to 130°C and 23 parts of N,N-dimethylbenzylamine are added. The reaction mixture is held at this temperature until the EEW has reached the level of 880 g/eq.

A mixture of 7097 parts of the crosslinking agent (B) [sic] and 90 parts of the additive K2000 (polyether, Byk Chemie/Germany) is then added and the resulting mixture is held at 100°C.

Half an hour later, 211 parts of butyl glycol and 1210 parts of isobutanol are added.

Immediately following this addition, a mixture of 467
5 parts of the precursor as per 2. (diethylenetriamine diketimine in methyl isobutyl ketone) and 520 parts of methylethanolamine is introduced into the reactor and the batch is brought to a temperature of 100°C. After a further half an hour, the temperature is raised to
10 105°C and 159 parts of N,N-diethylaminopropylamine are added.

75 minutes after the amine addition, 903 parts of Plastilit® 3060 (propylene glycol compound,
15 BASF/Germany) are added and the mixture is diluted with 522 parts of propylene glycol phenyl ether (mixture of 1-phenoxy-2-propanol and 2-phenoxy-1-propanol, BASF/Germany), in the course of which it is cooled rapidly to 95°C.

20 After 10 minutes, 14821 parts of the reaction mixture are transferred to a dispersing vessel. 474 parts of lactic acid (88% in water), dissolved in 7061 parts of deionized water, are added in portions with stirring. The mixture is subsequently homogenized for 20 minutes
25 before being diluted further with an additional 12600 parts of deionized water in small portions.

The volatile solvents are removed by vacuum distillation and then replaced by an equal volume of

deionized water.

The dispersion (A/B1) has the following characteristics:

5

Solids content: 33.8% (1 h at 130°C)

29.9% (1/2 h at 180°C)

Base content: 0.71 milliequivalents/g solids (130°C)

Acid content: 0.36 milliequivalents/g solids (130°C)

10 pH: 6.3

Particle size: 116 nm

(Mass average from photon correlation spectroscopy)

3.2 Preparation of the aqueous binder dispersion 15 (A/B2)

The preparation of the binder dispersion (A/B2) takes place in complete analogy to the binder dispersion (A/B1), except that directly following dilution with
20 propylene glycol phenyl ether 378 parts of K-KAT® XP 348 (bismuth 2-ethylhexanoate; 25% bismuth, King Industries, USA) are admixed to the organic stage with stirring. After cooling, 14821 parts of the reaction mixture are dispersed in exactly the same way as for
25 (A/B1).

The dispersion (A/B2) has the following characteristics:

Solids content: 33.9% (1 h at 130°C)

30.1% (1/2 h at 180°C)

Base content: 0.74 milliequivalents/g solids (130°C)

Acid content: 0.48 milliequivalents/g solids (130°C)

5 pH: 5.9

Particle size: 189 nm

(Mass average from photon correlation spectroscopy)

3.3 Preparation of the aqueous binder dispersion

10 (A/B3)

In a reactor equipped with a stirrer, reflux condenser, internal thermometer and inert gas inlet, 6824 parts of epoxy resin based on bisphenol A having an epoxy equivalent weight (EEW) of 188 are heated to 130°C under a nitrogen atmosphere together with 1984 parts of bisphenol A, 2527 parts of ethoxylated bisphenol A having an OH number of 222 (Dianol® 265, Akzo/Netherlands) and 597 parts of methyl isobutyl ketone. Then 16 parts of N,N-dimethylbenzylamine are added and the mixture is heated to 150°C and held at a temperature between 150 and 190°C for about 30 minutes. It is then cooled down to 140°C. This is followed by the addition of 21 parts of N,N-dimethylbenzylamine, and the temperature is maintained until the EEW has reached a level of 1120 g/eq.

Then 10113 parts of the crosslinking agent (B2) are added and the temperature is lowered to 100°C.

Subsequently, a mixture of 634 parts of the precursor (diethylenetriamine diketimine in methyl isobutyl ketone; cf. Section 2) and 597 parts of methylethanolamine are introduced into the reactor and
5 the reaction mixture is held at 115°C for one hour until a viscosity of approximately 6 dPa.s (50% dilution in methoxypropanol, cone and plate viscometer at 23°C). Then 648 parts of propylene glycol phenyl ether (mixture of 1-phenoxy-2-propanol and 2-phenoxy-1-
10 propanol, BASF/Germany) are added.

After 10 minutes the entire reaction mixture is transferred to a dispersing vessel. 609 parts of lactic acid (88% in water) and 152 parts of emulsifier mixture (mixture of 1 part of butyl glycol and 1 part of a
15 tertiary acetylene glycol (Surfynol 104, Air Products/USA)), dissolved in 30266 parts of deionized water, are added in portions with stirring.

The volatile solvents are removed by vacuum
20 distillation and then replaced by an equal volume of deionized water.

The dispersion (A/B3) has the following characteristics:

25

Solids content: 37.0% (1 h at 130°C)

34.1% (1/2 h at 180°C)

Base content: 0.53 milliequivalents/g solids (130°C)

Acid content: 0.32 milliequivalents/g solids (130°C)

pH: 6.6

Particle size: 150 nm

(Mass average from photon correlation spectroscopy)

5 **4. Preparation of aqueous solutions of polyvinyl
alcohol (co)polymers (D)**

**4.1 Preparation of an aqueous solution of poly(vinyl
alcohol-co-vinyl acetate) (D1)**

10

Poly(vinyl alcohol-co-vinyl acetate):

Mowiol® 47-88, Clariant/Germany

Weight average molar mass: 228 000 daltons (*)

15 Polyvinyl alcohol content: 89.2%

Polyvinyl acetate content: 10.8% (**)

(*) Weight average molar mass by light scattering
(+15% error) following reacetylation: 5 g of
poly(vinyl alcohol-co-vinyl acetate) are heated at
20 100°C for 24 hours with 75 ml of reacetylating
agent (pyridine/acetic anhydride/acetic acid =
1:10:10); reprecipitation from methanol in water.

(**) Calculated from ester number in accordance with
DIN 53401

25

A reactor equipped with a stirrer, reflux condenser,
internal thermometer and inert gas inlet is charged
with 28491 parts of deionized water at room
temperature. 1500 parts of poly(vinyl alcohol-co-vinyl

acetate) in the form of fine granules are stirred continuously into the initial water charge and the mixture is then heated to 80°C with stirring.

On reaching 80°C, the mixture is held with stirring for
5 two hours, the polymer being fully dissolved. This is followed by cooling to 35°C.

The viscous solution is stabilized against bacterial infestation with 9 parts of Parmetol® K40 (Schülke and
10 Mayr/Germany).

The solids content of the solution is 5.0% (1 h at 130°C).

4.2 Preparation of an aqueous solution of poly(vinyl 15 alcohol-co-vinyl acetate-co-ethylene) (D2)

Poly(vinyl acetate-co-ethylene):

Laboratory product, BASF AG, Germany

20 Weight average molar mass: 239 000 daltons (*)

Polyvinyl acetate content: 92.8% (**)

Polyethylene content: 7.2%

(*) Weight average molar mass by light scattering
(+15% error).

25 (**) Calculated from ester number in accordance with
DIN 53401

In a reactor equipped with a stirrer, reflux condenser and internal thermometer, 1000 ml of 1% strength

methanolic sodium hydroxide solution are heated to 50°C. Over the course of 30 minutes a solution of poly(ethylene-co-vinyl acetate) in methanol (300 g in 2000 ml of methanol) is added dropwise with stirring.

- 5 After the end of the addition, reaction is continued for 30 minutes and the precipitate is isolated, washed alkali-free with methanol and dried in vacuo at approximately 40°C.

- 10 The product formed was characterized:

Poly(vinyl alcohol-co-vinyl acetate-co-ethylene):

Weight average molar mass: 215 000 daltons (*)

- 15 Polyvinyl alcohol content: 83.3%

Polyvinyl acetate content: 9.5% (**)

Polyethylene content: 7.2%

- (*) Weight average molar mass by light scattering (+15% error) following reacetylation: 5 g of
20 poly(vinyl alcohol-co-vinyl acetate-co-ethylene) are heated at 100°C for 24 hours with 75 ml of reacetylating agent (pyridine/acetic anhydride/acetic acid = 1:10:10); reprecipitation from methanol in water.

- 25 (**) Calculated from ester number in accordance with DIN 53401

In analogy to the procedure in Section 4.1, an aqueous solution of poly(vinyl alcohol-co-vinyl acetate-co-

ethylene) is prepared.

The solids content of the solution is 5.0% (1 h at 130°C).

5 .

5. Preparation of the pigment pastes

5.1 Preparation of the gray pigment paste (P1)

10 5.1.1 Preparation of a grinding resin solution having tertiary ammonium groups

In accordance with EP 0 505 445 B1, Example 1.3, an aqueous-organic grinding resin solution is prepared by reacting, in the first stage, 2598 parts of bisphenol A diglycidyl ether (epoxy equivalent weight (EEW) 188 g/eq), 787 parts of bisphenol A, 603 parts of dodecylphenol and 206 parts of butyl glycol in a stainless steel reaction vessel in the presence of 4 parts of triphenylphosphine at 130°C until an EEW of 865 g/eq is reached. In the course of cooling, the batch is diluted with 849 parts of butyl glycol and 1534 parts of D.E.R.® 732 (polypropylene glycol diglycidyl ether, DOW Chemical, USA) and is reacted further at 90°C with 266 parts of 2,2'-aminoethoxyethanol and 212 parts of N,N-dimethylaminopropylamine. After 2 hours the viscosity of the resin solution is constant (5.3 dPa.s; 40% in Solvenon® PM (methoxypropanol, BASF/Germany); cone and plate

viscometer at 23°C). It is diluted with 1512 parts of butyl glycol and the base groups are partly neutralized with 201 parts of glacial acetic acid, and the product is diluted further with 1228 parts of deionized water and discharged.

This gives a 60% strength aqueous-organic resin solution whose 10% dilution has a pH of 6.0.

10 The resin solution is used in direct form for paste preparation.

5.1.2 Preparation of the pigment paste

15 For this purpose, a premix is first formed from 1897 parts of water and 1750 parts of the resin solution described above. Then 21 parts of Disperbyk® 110 (Byk-Chemie GmbH/Germany), 14 parts of Lanco Wax® PE W 1555 (Langer & Co./Germany), 42 parts of carbon black, 420 parts of aluminum hydrosilicate ASP 200 (Langer & Co./Germany), 2667 parts of titanium dioxide TI-PURE® R 900 (DuPont, USA) and 189 parts of di-n-butyltin oxide are added. The mixture is predispersed for 30 minutes under a high-speed dissolver stirrer. The mixture is subsequently dispersed in a small laboratory mill (Motor Mini Mill, Eiger Engineering Ltd, Great Britain) for from 1 to 1.5 h to a Hegmann fineness of less than or equal to 12 µm and adjusted to solids content with additional water.

A separation-stable pigment paste P1 is obtained.

Solids content: 60.0% (1/2 h at 180°C)

5 5.2 Preparation of the gray pigment paste (P2)

5.2.1 Preparation of a grinding resin solution having sulfonium groups

10 An aqueous-organic sulfonium grinding resin solution is prepared by reacting, in the first stage, 2632 parts of bisphenol A diglycidyl ether (epoxy equivalent weight (EEW) 188 g/eq), 985 parts of bisphenol A, and 95 parts of nonylphenol in a stainless steel reaction vessel in
15 the presence of 1 part of triphenolphosphine at 130°C until an EEW of 760 g/eq is reached. In the course of cooling, the temperature is lowered to 80°C with 996 parts of 2-butoxypropanol.

20 Then 603 parts of thiodiethanol (50% in water) are added and the mixture is stirred for 15 minutes. Following the addition of 661 parts of dimethylolpropionic acid and 152 parts of deionized water, the acid number is measured.

25

The reaction is over when the acid number is less than 5 (mg KOH per g solids). Then 10541 parts of deionized water are added in stages.

This gives a 28% strength aqueous-organic resin solution (solids content at 130°C, 60 min: 28.0%).

The resin solution is used in direct form for paste
5 preparation.

5.2.2 Preparation of a grinding resin solution having quaternary ammonium groups

10 First of all, in a reactor, 7507 parts of the 2-ethyl-
hexanol monourethane of tolylene diisocyanate (90%) are
added to 2040 parts of dimethylethanolamine at a rate
such that the temperature does not exceed 70°C. This
mixture is then diluted with 2199 parts of butyl
15 glycol, and 2751 parts of lactic acid (88%) and 2170
parts of deionized water are added. The temperature
rises to 90°C. After 3 hours the reaction product, used
subsequently as quaternizing reagent, is discharged.

20 An aqueous-organic grinding resin solution having
quaternary ammonium groups is prepared by reacting, in
the first stage, 3512 parts of bisphenol A diglycidyl
ether (epoxy equivalent weight (EEW) 188 g/eq), 1365
parts of bisphenol A, and 128 parts of xylene at 130°C
25 in a stainless steel reaction vessel in the presence of
4 parts of triphenylphosphine until an EEW of 740 g/eq
is reached. During the reaction the temperature is
raised to 180°C. The mixture is cooled and, at 125°C,
1947 parts of the 2-ethylhexanol monourethane of

tolylene diisocyanate (90%) are added. The temperature is held for about 2 hours until isocyanate groups are no longer detectable by IR. Following dilution with 4893 parts of butyl glycol, the temperature is adjusted
5 to 75°C and 3198 parts of the quaternizing reagent described above are added.

When the acid number is less than 1 (mg KOH per g solids), dilution is carried out with 1457 parts of
10 butyl glycol.

This gives a 56% strength resin solution (solids content at 130°C, 60 min: 56.0%).

15 The resin solution is used in direct form for paste preparation.

5.2.3 Preparation of the pigment paste

20 For this purpose, a premix is first formed from 1863 parts of water and 4119 parts of the above-described grinding resin solution having sulfonium groups (Section 5.2.1) and 422 parts of the above grinding resin solution having quaternary ammonium groups
25 (Section 5.2.2). Then 728 parts of aluminum hydrosilicate ASP 200 (Langer & Co./Germany), 185 parts of carbon black, 6142 parts of titanium dioxide TI-PURE® R 900 (DuPont, USA) and 3639 parts of di-n-butyltin oxide are added. The mixture is predispersed

for 30 minutes under a high-speed dissolver stirrer. The mixture is then dispersed in a small laboratory mill (Motor Mini Mill, Eiger Engineering Ltd, Great Britain) for from 1 to 1.5 h to a Hegmann fineness of
5 less than or equal to $12\ \mu\text{m}$ and is adjusted to solids with additional water.

A separation-stable pigment paste (P2) is obtained.

Solids content: 61.5% (1/2 h at 180°C)

10

6. Preparation of inventive electrocoat materials

The fractions of the components in the electrodeposition baths are set out in Tabs. 1, 2, and
15 3. Pigment-free and pigmented electrodeposition baths are the result.

These electrocoat materials comprise mixtures of in each case one aqueous dispersion (A/B) and deionized
20 water. In the selective cases, pigment paste (P) is added with stirring to the resultant mixtures.

The aqueous solutions of polyvinyl alcohol (co)polymers (D) may be incorporated by adding them to the binder
25 dispersion (A/B) or pigment paste (P) with stirring, or by subsequent addition to the binder/paste mixture, as in the present case.

Tab. 1:

Gray pigmented electrocoat materials based on the binder dispersion (A/B1) and the pigment paste (P1)

Electrocoat material	Comparative trial C1	Example 1	Example 2
Polyvinyl alcohol (co)polymer 1)	0 ppm 1)	600 ppm 1)	600 ppm 1)
Weight fractions (parts)			
Binder disp. (A/B1)	491	491	491
Pigment paste (P1)	120	120	120
Deionized water	389	377	377
Soln. of the polyvinyl alcohol (co)polymer (D1)		12	
(D2)			12
TOTAL	1000	1000	1000

5

1) Amount of polyvinyl alcohol (co)polymer (D) in the electrodeposition bath in ppm based on mass of electrodeposition bath

10 **Tab. 2:**

Unpigmented electrocoat (clearcoat) materials based on the binder dispersion (A/B2)

Electrocoat material	Comparative trial C2	Example 3	Example 4
Polyvinyl alcohol (co)polymer 1)	0 ppm 1)	1500 ppm 1)	600 ppm 1)
Weight fractions (parts)			
Binder disp. (A/B2)	498	498	498
Deionized water	502	462	462
Soln. of the polyvinyl alcohol (co)polymer (D1)		40	
(D2)			40
TOTAL	1000	1000	1000

1) Amount of polyvinyl alcohol (co)polymer in the electrodeposition bath in ppm based on mass of electrodeposition bath

Tab. 3:

Gray pigmented electrocoat materials based on the binder dispersion (A/B3) and the pigment paste (P2)

10

Electrocoat material	Comparative trial C3	Example 5	Example 6
Polyvinyl alcohol (co)polymer 1)	0 ppm 1)	600 ppm 1)	600 ppm 1)
Weight fractions			

(parts)

Binder disp. (A/B3)	416	416	416
Pigment paste (P2)	105	105	105
Deionized water	479	467	467
Soln. of the polyvinyl alcohol (co)polymer (D1)		12	
(D2)			12
TOTAL	1000	1000	1000

- 1) Amount of polyvinyl alcohol (co)polymer (D) in the electrodeposition bath in ppm based on mass of electrodeposition bath

5

7. Deposition of inventive electrocoat materials

After aging at room temperature for 5 days, deposition is carried out on a steel test panel connected as the cathode, with a series resistance of 150 ohm.

Water-rinsed, zinc-phosphated steel test panels from Chemetall 3) (Bo26 W OC) were used for this purpose. The deposition time was 2 minutes at a bath temperature of 32°C. The deposition voltage was chosen so as to give a baked film thickness of approximately 20 μm .

The deposited coating film is rinsed with deionized water and baked at 180°C for 20 minutes. The resultant

baked coating films were tested.

The test results may be seen from Tables 4 and 5.

5 7.1 Result of the depositions

As comparative examples, cathodically depositable electrodeposition baths without additions of polyvinyl alcohol (co)polymers were deposited (see also Section 10 6, Tabs. 1-3).

The film thicknesses reported are dry film thicknesses.

Tab. 4:

15 **Test results of electrodeposition baths based on the binder dispersions (A/B1) and (A/B2)*) with and without pigment paste (P1)**

*) Note: (A/B2) corresponds to (A/B1) with the addition 20 of bismuth carboxylate catalyst (see Section 3.2)

Electro- deposition baths	Gray		Unpigmented			
	pigmented		(clearcoat)			
Examples (see Sect. 6; Tabs. 1 + 2)	C1	1	2	C2	3	4
Binder dispersion	(A/B1)	"	"	(A/B2)*)	"	"

Pigment paste	(P1)	"	"	-	-	-
PVA1-CP						
solution (1)	-	(D1)	(D2)	-	(D1)	(D2)

Amount of PVA1-
CP in bath (2),

ppm	0	600	600	0	2000	2000
-----	---	-----	-----	---	------	------

Deposition on
zinc phosphate
steel test
panels (3)

Film thickness

(μ m)	20.7	20.9	20.2	20.6	19.2	19.3
Voltage, V	300	310	300	320	320	320

Electrical

rating (4) as

measure of edge

coverage, %	6	100	97	8	88	73
-------------	---	-----	----	---	----	----

Leveling (5)	2	3	3	2	2	3
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Corrosion

protection of

the 10 cycles

of alternating-

climate testing

(6)

Scribe creep,

105021 195007

Corrosion protection according to edge coating test Ford test method BI 127-01 (10)								-
No. of rust spots on a blade (10)	>80	19	21	>80		29		35
Oil splash compatibility (11) by BASF test method MEB0123A								
Cratered area as a % of total area (11):	>80	≤10	≤10	>80		≤10		≤10

Test results of electrodeposition baths based on the binder dispersions (A/B3) with pigment paste (P2)

Electrodeposition baths	Gray pigmented		
Examples (see Sect. 6; Tab. 3)	C3	5	6
Binder dispersion	(A/B3)	"	"
Pigment paste	(P2)	"	"
PVA1-CP solution (1)	-	(D1)	(D2)
Amount of PVA1-CP in bath (2), ppm	0	600	600
Deposition on zinc phosphate steel test panels (3)			
Film thickness (μm)	20.2	19.9	20.3
Voltage, V	310	310	310
Electrical rating (4) as measure of edge coverage, %			
	12	99	95
Leveling (5)	2	3	3
Corrosion protection of the 10 cycles of alternating-climate testing (6)			
Scribe creep, mm (7)	2.6	2.4	2.4
Surface rust (8)	1	1	1
Edge rust (9)	3	1	1
Corrosion protection according to edge coating test Ford test method BI 127-01 (10)			

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No. of rust spots on a blade (10) >80 22 24

Oil splash compatibility (11) by

BASF test method MEB0123A

Cratered area as a % of total area

(11): >80 ≤10 ≤10

(1) PVA1-CP: Polyvinyl alcohol copolymer

PVA1-CP solution: Polyvinyl alcohol copolymer
solution (for preparation see Section 4)

5

(2) PVA1 copolymer (polyvinyl alcohol copolymer):

Amount as solids based on electrodeposition bath
in ppm (see also Section 6, Tabs. 1 and 2)

10 (3) 2 minutes' deposition at 32°C on Bo26 W60 OC steel
test panels (water-rinsed, zinc-phosphated steel
test panels; water rinse pH = 6.0; Chemetall)

15 (4) This rating is determined by applying a voltage of
50-1000 V to the coated edge and determining the
insulating effect against breakdown. The test
panels used are again water-rinsed, zinc-
phosphated steel test panels (3) and are measured
on the 90° edge. The higher the electrical rating
20 (max. 100), the higher the insulating effect. The
higher the insulating effect, the better the
coating of the edge with a film of electrocoat
material.

T05021 "T9T600T"

- (5) 1 = best score; 5 = worst score
- (6) 10 cycles of alternating-climate testing in
5 accordance with VDA [German Automakers'
Association]
- (7) Creep [mm] = (total creep [mm] - scribe thickness
[mm]): 2
- 10
- (8) 0 = best score; 5 = worst score
- (9) 0 = best score; 5 = worst score
- 15 (10) Coated phosphated knife blades with a special 38°
cutting geometry (Embee blade No. 172; Embee
Corp., USA) are subjected to a 168 hour salt spray
test (Ford test method BI 103-01), after which the
number of rust spots appearing on the knife edge
20 is assessed. The smaller the number of rust spots,
the better the edge protection.
- (11) Oil splash compatibility test method MEB0123A of
BASF Coatings AG; test oil: Anticorit® RP 4107S
25 (Fuchs Mineralölwerke GmbH/Germany): the oil
splash compatibility of an electrocoat material is
investigated following contamination with a
crater-causing test oil during baking. The
percentage fraction of the cratered area is

evaluated. The smaller this area, the better the oil splash compatibility of the material. For the test, coated metal sample panels with unbaked air-dried electrocoat films are baked at 180°C for 15 minutes in the presence of a test oil/water mixture. The arrangement is such that the test oil splashes in a defined manner onto the sample panel during baking. As a result of this procedure, craters are formed in the baked coating, with the percentage area affected relative to the total area serving as a measure of the oil splash compatibility. For evaluation, the fraction of the cratered and uncratered area units is determined within a lattice network of defined lattice spacings. If, for example, max. 10% of the total area is cratered, the result is evaluated as <10%. The gradations are as follows: less than/equal to 10%, 11-20%, 21-40%, 41-80%, greater than 80%.

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What is claimed is:

1. The use of a water-soluble polyvinyl alcohol
(co)polymer or of a mixture of polyvinyl alcohol
5 (co)polymers as additives in aqueous
electrodeposition baths.
2. The use as claimed in claim 1, wherein the
polyvinyl alcohol (co)polymer is a copolymer of
10 vinyl alcohol and ethylenically unsaturated
monomers, preferably one ethylenically unsaturated
monomer or two or more ethylenically unsaturated
monomers, especially vinyl acetate, vinyl acetal,
ethylene and/or propylene.
- 15 3. The use as claimed in claim 1 or 2, wherein the
polyvinyl alcohol (co)polymer has a vinyl alcohol
fraction of from 50 to 99.9, preferably from 60 to
99.9, with particular preference from 70 to 99,
20 and in particular from 80 to 99 mol%.
4. The use as claimed in any of claims 1 to 3,
wherein the weight average molecular mass of the
polyvinyl alcohol (co)polymer is from 10 000 to
25 500 000, preferably from 15 000 to 320 000, and in
particular from 20 000 to 300 000 daltons.
5. The use as claimed in any of claims 1 to 4,

wherein the fraction of polyvinyl alcohol (co)polymer in the electrodeposition bath is from 2 to 10 000 ppm, preferably from 20 to 5000 ppm, and in particular from 300 to 1500 ppm, based in each case on the total weight of the electrodeposition bath.

6. An aqueous electrodeposition bath comprising
 - (A) a cathodically or anodically depositable binder,
 - (B) optionally a crosslinking agent,
 - (C) optionally, customary coatings additives,
 - (D) a dissolved polyvinyl alcohol (co)polymer as set forth in any of claims 2 to 5.
7. A method of coating electrically conductive substrates, in which
 - (1) the electrically conductive substrate is dipped into an electrodeposition bath as claimed in claim 6,
 - (2) the substrate is connected as the cathode or anode,
 - (3) a film is deposited on the substrate by means of direct current,
 - (4) the coated substrate is removed from the electrodeposition bath,
 - (5) the deposited coating film is baked, and,
 - (6) optionally, following step (5), a primer-surfacer, and/or a stonechip protectant

material and a solid-color topcoat material,
or alternatively a basecoat material and a
clearcoat material, are applied and baked,
the basecoat material and the clearcoat
5 material being applied and baked in
particular by the wet-on-wet technique.

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DECLARATION FOR UTILITY OR DESIGN PATENT APPLICATION (37 CFR 1.63) <input checked="" type="checkbox"/> Declaration Submitted with initial Filing or <input type="checkbox"/> Declaration Submitted after initial Filing (surcharge (37 CFR 1.16 (e)) required	Attorney Docket No.	IN - 5542
	First Named Inventor	Hardy REUTER
	COMPLETE IF KNOWN	
	Application Number	
	Filing Date	Wednesday, December 05, 2001
	Group Art Unit	
	Examiner Name	

As below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name.

I believe I am the original, first and sole inventor (if only name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention **entitled:**

ELECTRODEPOSITION BATH WITH WATER-SOLUBLE POLYVINYL ALCOHOL (CO)POLYMERS

(Title of the Invention)

The specification of which:

☒ is attached hereto

☒ Was filed on **29 June 2000** as United States Application or PCT International Application Number **PCT/EP00/06035** and was amended on _____ (if applicable).

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment specifically referred to above.

I acknowledge the duty to disclose information which is material to patentability as defined in 37 CFR 1.56, including for continuation-in-part applications, material information which became available between the filing date of the prior application and the national or PCT international filing date of the continuation-in-part application

I hereby claim foreign priority benefits under 35, U.S.C § 119(a)-(d) or 365(b) of any foreign application(s) for patent or inventor's certificate or 365 (a) of any PCT international application(s) designating at least one country other than the United States of America listed below and have also identified below, by checking the box, any foreign application(s) for patent or inventor's certificate, or any PCT international application(s) having a filing date before that of the application on which priority is claimed.

Prior Foreign Application Number(s)	Country	Foreign Filing Date DATE/MONTH/YEAR	Priority Not Claimed	Certified Copy Attached?	
				Yes	No
199 30 060.7	GERMANY	30. JUNE 1999	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
			<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Additional foreign application number are listed on a supplemental priority data sheet PTO/SB/02B attached hereto:

I hereby claim the benefit under 35 U.S.C 119(e) of any United States provisional application(s) listed below:

APPLICATION NUMBER(S)	FILING DATE	<input type="checkbox"/> Additional provisional application numbers are listed on a supplemental priority data sheet PTO/SB/02B attached hereto

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Name OF SOLE OR FIRST NVENTOR:



A petition has been filed for this unsigned inventor

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(first and middle [if any])

Hardy

Family Name
or Surname

REUTER

Inventor's
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16.11.01

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Name OF SECOND INVENTOR:

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A petition has been filed for this unsigned inventor

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Inventor's
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16. 11. '01

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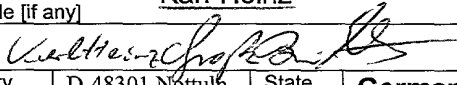
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Name of Additional Joint Inventor, If any: <input type="checkbox"/>				A petition has been filed for this unsigned inventor <input type="checkbox"/>			
Given Name (first and middle [if any]) <u>Karl-Heinz</u>				Family Name or Surname <u>GROSSE-BRINKHAUS</u>			
Inventor's Signature 				Date <u>16.11.01</u>			
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City	<u>D-48301 Nottuln</u>	State	<u>Germany</u>	Zip			

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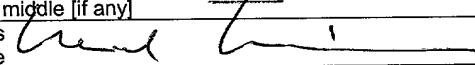
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Name of Additional Joint Inventor, If any: <input type="checkbox"/>				A petition has been filed for this unsigned inventor			
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Inventor's Signature 				Date <u>16.11.01</u>			
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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Name of Additional Joint Inventor, If any: ☐ A petition has been filed for this unsigned inventor

Given Name (first and middle [if any])		Walter		Family Name or Surname		JOUCK	
Inventor's Signature				Date		16.11.01	
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